1. (Previously Amended) A compound that is an oxazolidinone derivative of the formula (I)

or a salt thereof, or a stereoisomer thereof, where

 $R^1$  represents -NHR $^4$  wherein  $R^4$  represents thio( $C_1$ - $C_{10}$ )acyl, -C(=S)-cyclo( $C_3$ - $C_8$ )alkoxy, -C(=S)-( $C_1$ - $C_{10}$ )alkoxy, -C(=S)-( $C_2$ - $C_{10}$ )alkenyloxy, -C(=S)-aryloxy, -(C=S)-S-( $C_1$ - $C_{10}$ )alkyl, -(C=S)-NH-( $C_1$ - $C_1$ 0)alkyl, -C(=S)-N-(( $C_1$ - $C_1$ 0)alkyl)2, -C(=S)-NH-( $C_2$ - $C_1$ 0)alkenyl, (C=S)-( $C_1$ - $C_1$ 0)alkoxy, -(C=S)-(C=O)-aryloxy, -C(=S)-O-( $C_1$ - $C_1$ 0)alkyl, C(=S)-C(=S)-( $C_1$ - $C_1$ 0)alkyl, -C(=S)-C(=S)-aryl, -C(=S)-thiomorpholinyl or -C(=S)-pyrrolidinyl;  $R^2$  and  $R^3$ , which may be the same or different, are each independently hydrogen, halogen, ( $C_1$ - $C_1$ 0)alkyl, halogenated ( $C_1$ - $C_1$ 0)alkyl, cyano, nitro,  $SR^a$ ,  $NR^a$ , or  $OR^a$ , in which  $R^a$  is hydrogen, ( $C_1$ - $C_1$ 0)alkyl or halogenated ( $C_1$ - $C_1$ 0)alkyl;

 $Y^2$   $Y^3$  is a heterocyclic moiety in which is a 5-membered heterocyclic skeleton, Z represents =CH, -CH<sub>2</sub> or NR<sup>b</sup>, where R<sup>b</sup> is hydrogen or a moiety, which may be substituted or unsubstituted, straight chain or branched, selected from the group consisting of  $(C_1-C_{10})$ alkyl,  $(C_2-C_{10})$ alkenyl,  $(C_3-C_8)$ cycloalkyl, hydroxy $(C_1-C_{10})$ alkyl,  $(C_1-C_{10})$ alkylhydroxy,  $(C_1-C_{10})$ alkylamino, amino $(C_1-C_{10})$ alkyl,  $(C_1-C_{10})$ alkoxy, aryl, aralkyl, aryloxy,  $(C_1-C_{10})$ alkylcarbonyl, arylcarbonyl,  $(C_1-C_{10})$  alkoxycarbonyl and aryloxycarbonyl;

Y<sup>1</sup> represents =O or =S group and Y<sup>2</sup> and Y<sup>3</sup> independently represent hydrogen, halogen, cyano, nitro, formyl, hydroxy, amino, =O, =S group or substituted or

unsubstituted groups selected from  $(C_1-C_{10})$ alkyl, hydroxy $(C_1-C_{10})$ alkyl,  $(C_1-C_{10})$ alkylhydroxy,  $(C_1-C_{10})$ alkoxy $(C_1-C_{10})$ alkyl,  $(C_1-C_{10})$ alkyl,  $(C_1-C_{10})$ alkylcarbonyl, arylcarbonyl, carboxy $(C_1-C_{10})$ alkyl,  $(C_1-C_{10})$ alkylcarbony $(C_1-C_{10})$ alkyl, arylcarbonylamino $(C_1-C_{10})$ alkyl,  $(C_1-C_{10})$ alkylcarbonyloxy $(C_1-C_{10})$ alkyl, amino $(C_1-C_{10})$ alkyl, mono $(C_1-C_{10})$ alkylamino, di $(C_1-C_{10})$ alkylamino, arylamino,  $(C_1-C_{10})$ alkoxy, aryl, aryloxy, aralkyl, heteroaryl, heteroaralkyl, heterocyclyl or heterocycloalkyl;  $(C_1-C_1)$ alkylamino, arylamino arylamino, carbon atoms together may also form a substituted or unsubstituted 5 or 6 membered aromatic or non-aromatic cyclic structure, optionally containing one or two hetero atoms selected from oxygen, sulfur and nitrogen.

2. (Original) The compound of claim 1 having the structure

whrein X<sup>1</sup> is oxygen or sulfur.

3.. (Original) The compound of claim 1 having the structure

$$R^{b}-N$$
 $N$ 
 $R^{2}$ 
 $R^{2}$ 
 $R^{3}$ 
 $R^{1}$ 

wherein X<sup>1</sup> is oxygen or sulfur.

4. (Original) The compound of claim 1 having the structure

wherein  $X^1$  is oxygen or sulfur, and is a substituted or unsubstituted 5- or 6-membered aromatic or non-aromatic cyclic structure optionally having one or two hetero atoms, formed by  $Y^2$  and  $Y^3$ .

5-6. (Canceled).

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- 7. (Original) The compound of claim 4, wherein said cyclic structure formed by  $Y^2$  and  $Y^3$  is benzene, pyridine, pyrrolidine, furan thiophene, morpholine, piperazine or pyrrole.
- 8. (Previously Amended) The compound of formula (I) as defined according to claim 1 which is selected from:

9-78. (Canceled).

- 79. (Previously Amended) A pharmaceutical composition comprising a) an antibacterially effective amount of the compound of claim 1; and b) a pharmaceutically acceptable carrier.
- 80-88. (Canceled)
- 89. (Previously Presented) The compound of claim 1, having the structure

91. (Previously Presented) The compound of claim 1, having the structure

92. (Previously Presented) The compound of claim 1, having the structure

95. (Previously Presented) The compound of claim 1, having the structure

96. (Previously Presented) The compound of claim 1, having the structure

- 97. (Canceled).
- 98. (Previously Presented) The compound of claim 1, having the structure

101. (Previously Presented) The compound of claim 1, having the structure

102. (Previously Presented) The compound of claim 1, having the structure

103. (Previously Presented) The compound of claim 1, having the structure

106. (Previously Presented) The compound of claim 1, having the structure

107 - 108. (Canceled).

109. (Previously Presented) The compound of claim 1, having the structure

110. (Canceled).

111. (Previously Presented) The compound of claim 1, having the structure

113 - 114. (Canceled).

115. (Previously Presented) The compound of claim 1, having the structure

116. (Previously Presented) The compound of claim 1, having the structure

117. (Previously Presented) The compound of claim 1, having the structure